



IMUNON R&D Day: Opportunity to Hear Clinical Trial Investigators Discuss Significant Potential of IMNN-001 to Redefine Ovarian Cancer Treatment

November 7, 2025

Investors and stakeholders invited to attend in person in New York City or via live webcast on Monday, November 10th at 8:00 a.m. ET

LAWRENCEVILLE, N.J., Nov. 07, 2025 (GLOBE NEWSWIRE) -- IMUNON, Inc. (Nasdaq: IMNN), a clinical-stage company in Phase 3 development with its DNA-mediated immunotherapy, will host an R&D Day on Monday, November 10, 2025 at 8:00 a.m. ET in New York City featuring in-depth discussions with experts including principal investigators leading the Phase 3 OVATION 3 clinical trial and Phase 2 minimal residual disease (MRD) trial of IMNN-001, a novel IL-12 immunotherapy in development for the treatment of women with advanced ovarian cancer.

"This is an opportunity to hear directly from the investigators driving IMNN-001 clinical progress forward—physicians who see firsthand the urgent need for new ovarian cancer treatment options and the potential clinical impact of our novel IMNN-001 immunotherapy," said Stacy R. Lindborg, Ph.D., president and chief executive officer of IMUNON. "The data and insights shared will be consequential for understanding the future of ovarian cancer treatment and how we are leading the effort to bring the first advance in the standard of care that could transform frontline treatment, which has not seen innovation for about 30 years."

The event will include presentations from:

- Premal H. Thaker, M.D. (Washington University School of Medicine) – Unmet need and OVATION 2 trial data including survival benefits
- Amir A. Jazaeri, M.D. (MD Anderson Cancer Center) – Phase 2 MRD study results and immune activation mechanism overview
- Giorgio Paulon, Ph.D. (Berry Consultants, LLC) – Phase 3 trial statistical design and path to approval
- Douglas V. Faller, M.D., Ph.D. (IMUNON) – Phase 3 trial enrollment momentum and clinical milestones

To register to attend the event in person or virtually, please RSVP by clicking [here](#).

About IMUNON

IMUNON is a clinical-stage biotechnology company focused on advancing a portfolio of innovative treatments that harness the body's natural mechanisms to generate safe, effective and durable responses across a broad array of human diseases, constituting a differentiating approach from conventional therapies. IMUNON is developing its non-viral DNA technology across its modalities. The first modality, TheraPlas[®], is developed for the gene-based delivery of cytokines and other therapeutic proteins in the treatment of solid tumors where an immunological approach is deemed promising. The second modality, PlaCCine[®], is developed for the gene delivery of viral antigens that can elicit a strong immunological response.

The Company's lead clinical program, IMNN-001, is a DNA-based immunotherapy for the localized treatment of advanced ovarian cancer that has completed multiple clinical trials including one Phase 2 clinical trial (OVATION 2) and is currently conducting a Phase 3 clinical trial (OVATION 3). IMNN-001 works by instructing the body to produce safe and durable levels of powerful cancer-fighting molecules, such as interleukin-12 and interferon gamma, at the tumor site. Additionally, the Company has completed dosing in a first-in-human study of its COVID-19 booster vaccine (IMNN-101). The Company will continue to leverage these modalities and to advance, either directly or through partnership, the technological frontier of plasmid DNA to better serve patients with difficult-to-treat conditions. For more information, please visit www.imunon.com.

Forward-Looking Statements

IMUNON wishes to inform readers that forward-looking statements in this news release are made pursuant to the "safe harbor" provisions of the Private Securities Litigation Reform Act of 1995. All statements, other than statements of historical fact, including, but not limited to, statements regarding the timing of enrollment of the Company's clinical trials, the potential of any therapies developed by the Company to fulfill unmet medical needs, the market potential for the Company's products, if approved, the potential efficacy and safety profile of our product candidates, and the Company's plans and expectations with respect to its development programs more generally, are forward-looking statements. We generally identify forward-looking statements by using words such as "may," "will," "expect," "plan," "anticipate," "estimate," "intend" and similar expressions (as well as other words or expressions referencing future events, conditions or circumstances). Readers are cautioned that such forward-looking statements involve risks and uncertainties including, without limitation, uncertainties relating to unforeseen changes in the course of research and development activities and in clinical trials, including the fact that interim results are not necessarily indicative of final results; the uncertainties of and difficulties in analyzing interim clinical data; the significant expense, time and risk of failure in conducting clinical trials; the need for IMUNON to evaluate its future development plans; possible actions by customers, suppliers, competitors or regulatory authorities; and other risks detailed from time to time in IMUNON's filings with the Securities and Exchange Commission. IMUNON assumes no obligation, except to the extent required by law, to update or supplement forward-looking statements that become untrue because of subsequent events, new information or otherwise.

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